

State of Oklahoma Oklahoma Health Care Authority Xolair[®] (Omalizumab) Prior Authorization Form

| Member Name: | | er Name: | Date of Birth: | Member ID#: | | | |
|--|-------|---|--|--|--|--|--|
| | | | Drug Information | n | | | |
| | Ph | vsician billing (HCPCS code: | \ \ \ \ Pharm | nacy billing* (NDC: | | | |
| ☐ Physician billing (HCPCS code:_*If medication is being billed by a pharmacy, the medic | | | cation should be shipped to the heal | th care facility where it will be administered. | | | |
| Dos | se:_ | | Regimen: | Fill Date: | | | |
| | | | Billing Provider Inform | nation | | | |
| Soc | ner | rCare Provider ID: | Provider Na | nme: | | | |
| Provider Phone: | | | Provider Fa | x: | | | |
| Nan | ne c | of outpatient health care facili | ty where Xolair [®] will be d | elivered to and administered at: | | | |
| | | | | | | | |
| | | | Prescriber Informat | | | | |
| Prescriber NPI: | | | Prescriber Name: | | | | |
| Prescriber Phone: | | | Prescriber Fax: | Specialty: | | | |
| | | | Clinical Information | on | | | |
| | | mation must be provided and So 's drug history will be reviewed լ | | h further requested documentation. The | | | |
| Page | e 1 o | of 2—Please complete and return | all pages. Failure to compl | lete all pages will result in processing delays. | | | |
| For I | nitia | al Authorization: | | | | | |
| | | t is the diagnosis for which the med | lication is being prescribed? | | | | |
| | | Severe Persistent Asthma [as per National Asthma Education and Prevention Program guidelines] | | | | | |
| | | ☐ Chronic Idiopathic Urticaria | | | | | |
| | | Other, please list: | | | | | |
| | | iagnosis is Severe Persistent Asthma , please provide the following (Initial approvals will be for the duration of 12 months): | | | | | |
| A. Does member have a positive skin test to at least 1 perennial aeroallergen? Yes No | | | | | | | |
| | | i. If "Yes", please list perennial a | aeroallergen(s): | | | | |
| | | Please provide member's baseline | | | | | |
| | | Please provide member's weight: _ | | | | | |
| | | | • | g/day fluticasone propionate or equivalent daily dose | | | |
| | | | rears) used compliantly for at | least the past 3 months? Yes No | | | |
| | _ | i. Drug/Dose: | | | | | |
| | | | ealth care setting by a health | care professional prepared to manage | | | |
| | | anaphylaxis? Yes No | | | | | |
| | | | | or pulmonary specialist within the last 12 months (or | | | |
| | | • | a supervising physician who | is an allergist, pulmonologist, or pulmonary | | | |
| | | specialist)? Yes No | of appointing | | | | |
| | _ | i. If "Yes", please include name | | zations and/or ER visits in the past 12 | | | |
| | | months: | es or astrima related nospitali. | Zations and/or ER visits in the past 12 | | | |
| | | Is member dependent on systemic | corticosteroids to prevent se | rious asthma exacerbations? Yes No | | | |
| | | • | | ng (Initial approvals will be for the duration of 3 months): | | | |
| | | Have other forms of urticaria been | | .g (a. approvate tim bo for the daration of a monthla). | | | |
| | | Have other potential causes of urtic | | No | | | |
| | | Please provide member's Urticaria | | | | | |
| | | - | , , , | | | | |
| | | | Page 1 of 2 | | | | |

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy Pharmacy Management Consultants Product Based Prior Authorization Unit Fax: 1-800-224-4014

Phone: 1-800-522-0114 Option 4

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| Member Name: | Date of Birth: | Member ID#: |
|--|--|--|
| | Clinical Informati | on |
| Page 2 of 2—Please complete a | nd return <u>all</u> pages. <i>Failure to comբ</i> | olete all pages will result in processing delays. |
| D. Was Xolair [®] prescribed be supervising physician that i. If "Yes", please included in the supervision of | thic Urticaria, please provide the follow by an allergist, immunologist, or dermant at is an allergist, immunologist, or dermant and de name of specialist: of a second generation H ₁ antihistamin and 4 weeks? Yes No avide the medication used, dose prescri | tologist (or an advanced care practitioner with a natologist)? Yes No be dosed 4 times the maximum FDA dose within the bed, and dates of use: |
| 4-week trial is not a | ppropriate for this member: red in a health care setting by a health | Dates of use: us less than 4 weeks, please provide a reason why a use care professional prepared to manage |
| Urticaria Activity Score (UAS) a. If there has been no imp | es Chronic Idiopathic Urticaria, please p : Date assessed: rovement in member's UAS score, plea | provide member's current ase provide additional clinical information to support |
| Additional Information: | | |
| | | |
| Compliance with all of the prior | authorization criteria is a condition | for payment for this drug by SoonerCare. All |
| information must be provided a drug history will be reviewed pr | | further requested documentation. The member's |
| Prescriber Signature: | | Date:ccurate and verifiable in patient records.) |
| (By signature, the physician confir | ms the criteria information above is ac | curate and verifiable in patient records.) |
| Pharmacist Signature: | | Date: |
| Pease do not send in chart note | es. Specific information/documentat | tion will be requested if necessary. Failure to |

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